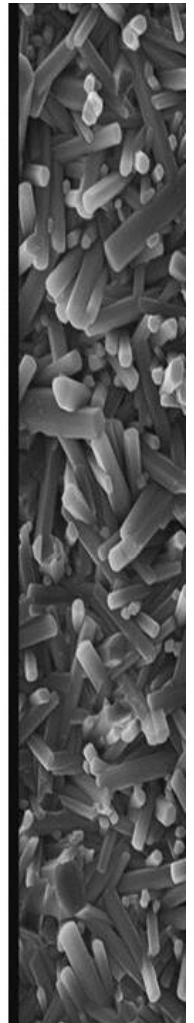


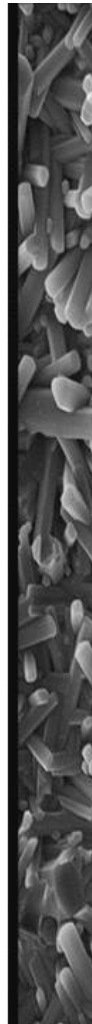


Amedica Corporation (NASDAQ:AMDA)
Corporate Presentation | June 2016



Free Writing Prospectus

Amedica Corporation (“Amedica”, “we”, “our”, “us”, or the “Company”) is providing this presentation, which highlights basic information about us and the offering to which this communication relates. Because it is a summary, it does not contain all of the information that you should consider before investing in our securities. We have filed a registration statement, including a prospectus, with the U.S. Securities and Exchange Commission (“SEC”) for the offering to which this communication relates. The registration statement has not yet become effective. Before you invest, you should read the prospectus in the registration statement, including the risk factors described therein, and other documents we have filed with the SEC for more complete information about us and the offering. You may access these documents for free by visiting EDGAR on the SEC website at www.sec.gov. Alternatively, the Company or any underwriter or dealer participating in the offering will arrange to send you the prospectus and/or any supplements thereto if you contact Ladenburg Thalmann & Co. Inc., Attention: Prospectus Department, 570 Lexington Ave New York, NY, 10022 email: prospectus@ladenburg.com.



Forward Looking Statements

This presentation contains forward-looking information that is based on beliefs and assumptions of Amedica's management and on information currently available to us. These forward-looking statements involve significant risks and uncertainties, including those discussed in this presentation and others that can be found in the "Risk Factors" section of the Registration Statement on Form S-1 of Amedica Corporation, filed with the SEC on June 8, 2016. All statements, other than statements of historical facts, included in this presentation regarding our strategy, products, future operations, projected expenses, products' placements, performance and acceptance, prospects, plans, management's objectives, and the growth of the overall market for our products in general and certain products in particular are forward looking statements.

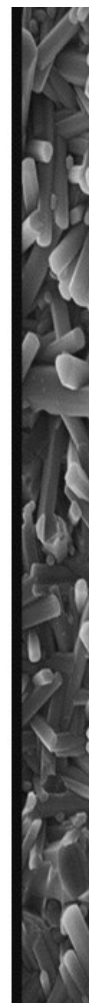
These forward-looking statements relate to future events or to our future financial performance and involve known and unknown risks, uncertainties, and other factors that may cause Amedica's or our industry's actual results, levels of activity, performance or achievements to be materially different from any future results, levels of activity, performance or achievements expressed or implied by these forward-looking statements. You should not rely on forward-looking statements as predictions of future events. We are unable to give any assurances concerning actual future revenues or that the properties of any of our product candidates will be sufficient to support U.S. Food and Drug Administration or other regulatory clearance or approval. Amedica is providing the information in this presentation as of this date and does not undertake to update any forward-looking information contained in this presentation as a result of new information, future events or otherwise.



Transaction Overview

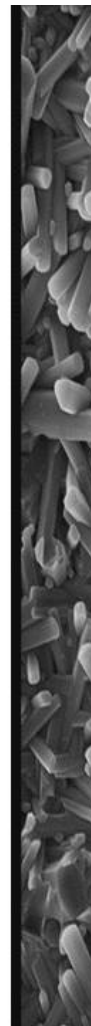
Issuer:	<ul style="list-style-type: none"> Amedica Corporation
Exchange:	<ul style="list-style-type: none"> NASDAQ: AMDA
Offering Size:	<ul style="list-style-type: none"> \$11.5 million
Securities Offered:	<ul style="list-style-type: none"> Class A Units consisting of common stock, Series E Warrants exercisable for shares of common stock and Series F Warrants exercisable for shares of common stock; or Class B Units consisting of Series A Convertible Preferred Stock, Series E Warrants exercisable for shares of common stock and Series F Warrants exercisable for shares of common stock¹
Use of Proceeds:	<ul style="list-style-type: none"> Continue to build sales, marketing and distribution capabilities Clinical development of pipeline products General Corporate Purposes Redeem senior convertible note held by MG Partners II Redeem subordinated convertible promissory note held by Riverside Merchant Partners, LLC To support debt service under existing senior secured credit facility with Hercules Technology Group
Joint Book-runners:	<ul style="list-style-type: none"> Ladenburg Thalmann & Co. Inc. Maxim Group LLC

1. Class B Units are to be offered to purchasers whose purchase of Class A Units in this offering would result in the purchaser together with its affiliates and certain related parties, beneficially owning more than 4.99% (or, at the election of the purchaser 9.99%) of the Company's common stock following the consummation of the offering.



About Us

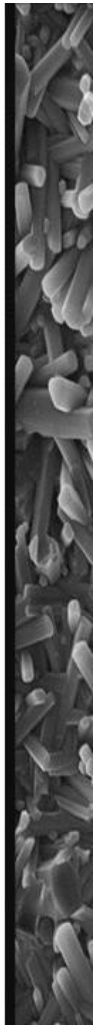
Biomaterial	Silicon nitride (Si_3N_4) is an advanced ceramic	<ul style="list-style-type: none">• Proven bone growth and anti-infective properties• Superior imaging, strength, and wear resistance
Hybrid Commercial Strategy	Traditional distribution with private label and OEM partnerships	<ul style="list-style-type: none">• Growth through expanded surgeon and distribution relationships• Validates Si_3N_4 benefits <u>AND</u> penetrates market w/ improved operating margins
Broad Product Portfolio	Interbody fusion devices and Metals business with pull-through effect of Si_3N_4	<ul style="list-style-type: none">• >25,000 Si_3N_4 spinal fusion devices implanted• Metals business includes facet and pedicle screw systems
Manufacturing	In-house manufacturing and Kyocera as secondary supplier	<ul style="list-style-type: none">• Only FDA & CE cleared Si_3N_4 medical device manufacturing facility• Quick in-house prototyping and development• Kyocera partnership allows for improved margins & rapid expansion
Seasoned Management	Surgery, science, manufacturing, & product development expertise	<ul style="list-style-type: none">• Superb people behind the product



Upcoming Milestones

2016 Milestones

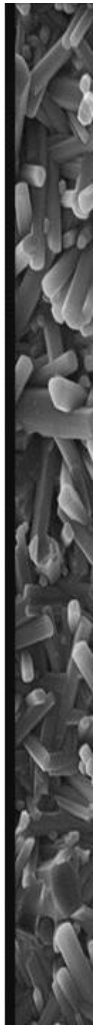
- Further reductions in operational cash burn
- Final determination by FDA for composite silicon nitride fusion device
- Report 12-month SNAP clinical data
- Sign four additional OEM or private label agreements
- Continue robust scientific publication strategy
- Additional product launches
 - Next generation of silicon nitride
 - New pedicle screw system
 - Expanded lateral lumbar implant sizes



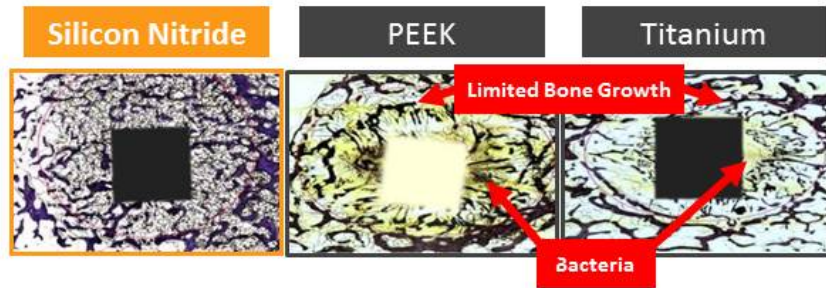
Silicon Nitride: The Ideal Biomaterial

	Silicon Nitride	Plastic	Allograft	Metals
Antibacterial	✓	✗	✗	✗
Bone Growth	✓	✗	✓	✗
Imaging Capability	✓	~	✓	✗
Strength, Hardness, Fracture Resistance	✓	✗	✗	✓
Bio Compatible	✓	✗	✓	~

Silicon nitride **actively participates** in the patient healing process



Anti-Infective & Osteopromotive



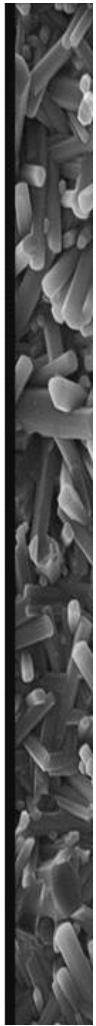
Bacteria growth after implant

	Silicon Nitride	PEEK	Titanium
On implant	0%	95%	67%
Growth <i>in situ</i>	0%	88%	21%

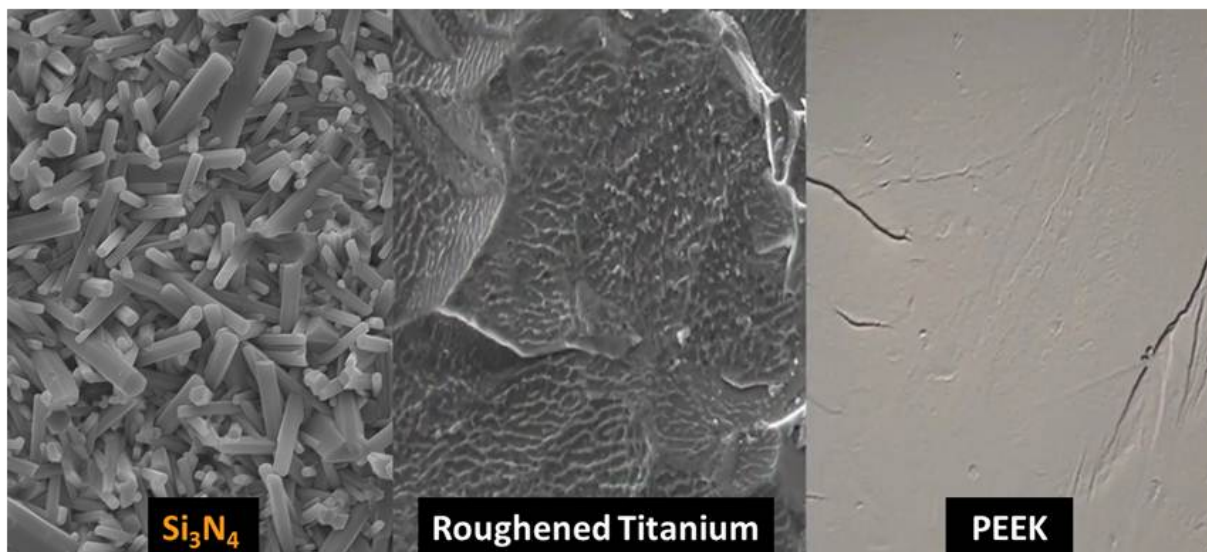
Bone growth in presence of bacteria

	Silicon Nitride	PEEK	Titanium
On implant	23%	5%	9%
Growth <i>in situ</i>	41%	21%	26%

In vivo Wistar Rat Calvarial Study: Histology at 3 months after implantation with inoculation *S. epidermis*



Material Surface Topography Makes the Difference



Silicon nitride has **superior surface area** for an optimal bone friendly environment

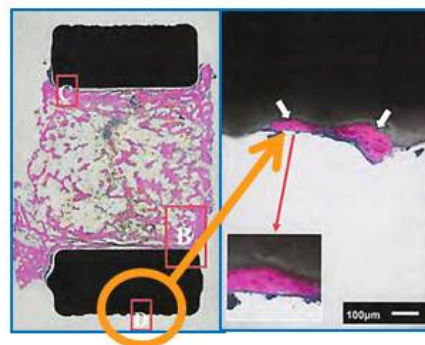
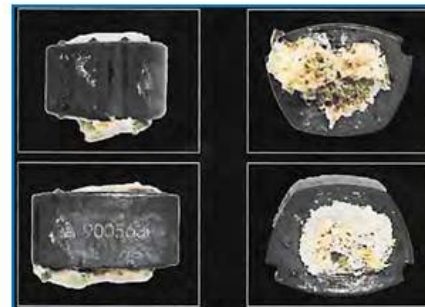
Case Study¹ - Silicon Nitride Encourages Bone Fusion

Details

- 58-year-old male
- Silicon nitride cervical implant C3-C4
- Length of implantation: 10 months
- Revision due to anatomical change requiring cervical plate removal

Histology

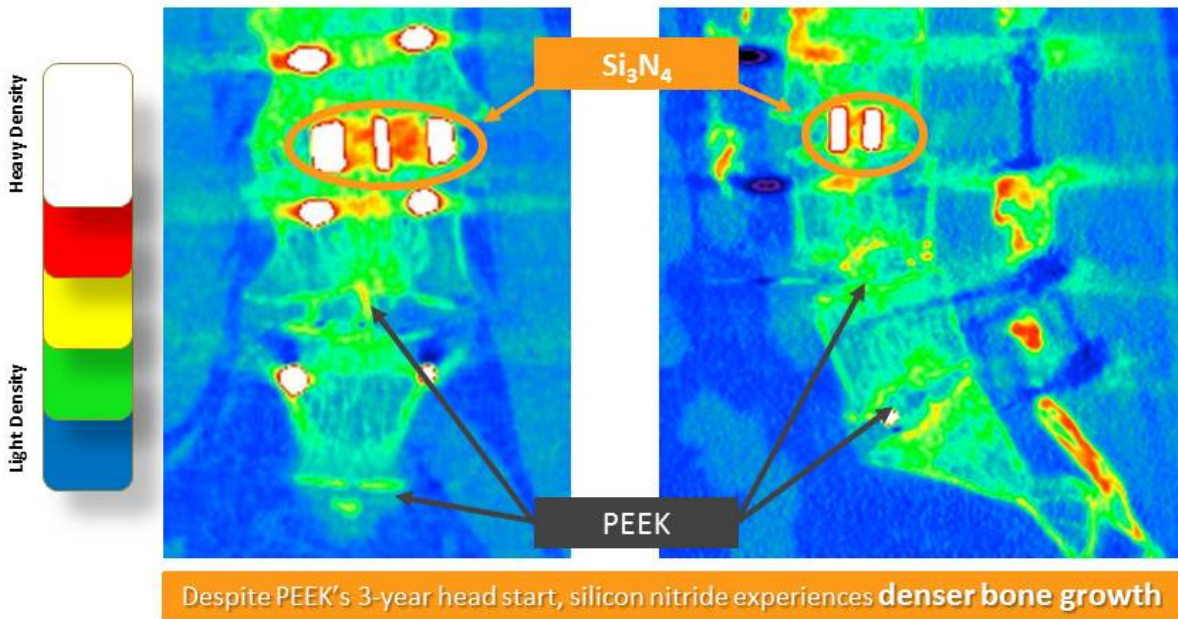
- Mature, healthy, viable bone throughout the graft hole area
- Good connectivity between vertebrae
- **Appositional bone index of 19.1% compared to 1.3% typical for PEEK²**



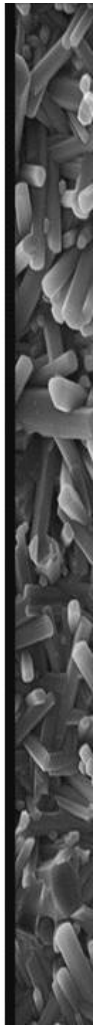
¹ R. Bloebaum, University of Utah BJRL Report, AS 86/13, (2013).

² Sinclair, Sarina K., et al. "Host bone response to polyetheretherketone versus porous tantalum implants for cervical spinal fusion in a goat model." *Spine* 37.10 (2012): E571-E580.

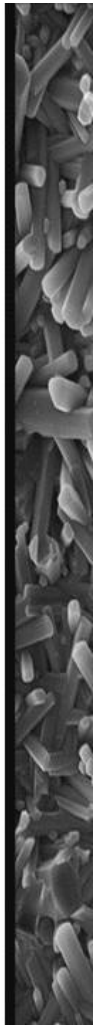
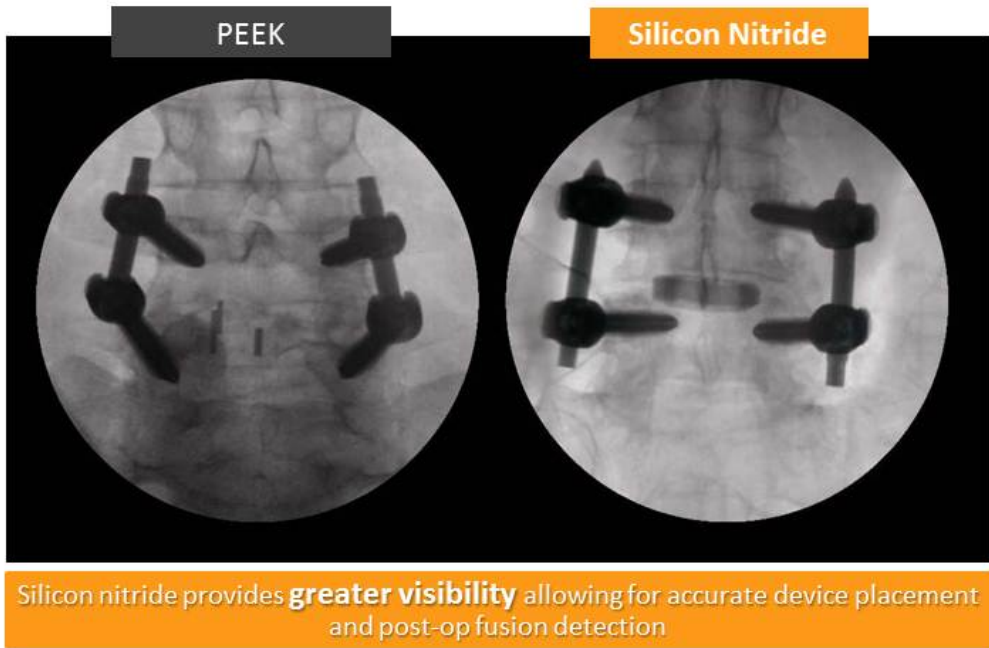
Case Study¹ - Silicon Nitride vs. PEEK in Same Patient



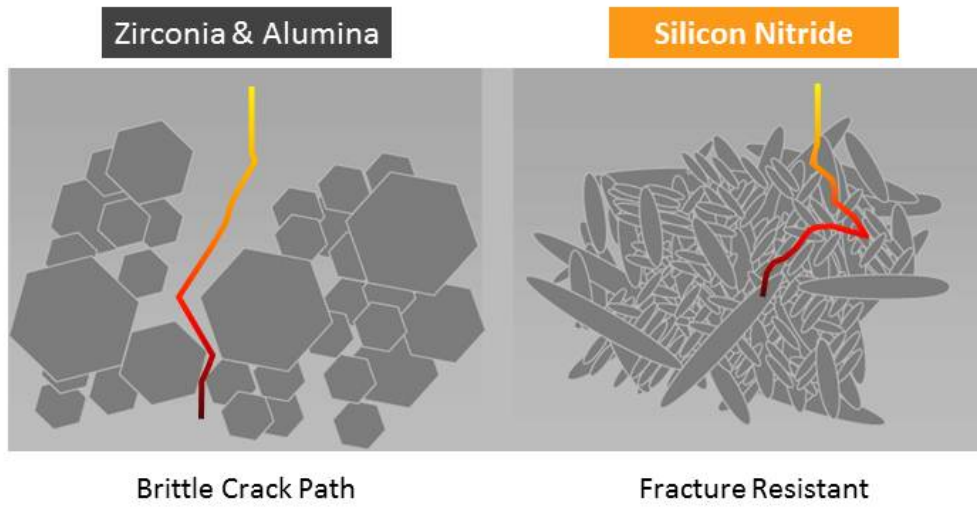
¹Muhanna, Nabil, MD, "A Retrospective Radiographic Review of PEEK and Silicon Nitride Spinal Implants in the Same Patient"



Easier to see...

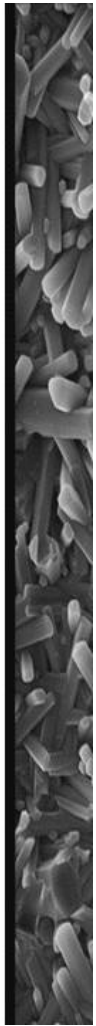


Strength & Reliability from Crack Resistance



Superior burst strength and fracture toughness vs. zirconia & alumina

18.S. Bai, M.N. Rehaman / Acta Biomaterialia 8 (2012) 2889–2898



Proprietary Silicon Nitride Types



Solid: As-Fired and Polished

As-fired promotes bone growth
Polished used for articulating applications

Porous: Cancellous (CsC)

Biologic substitute allowing for bone in-growth

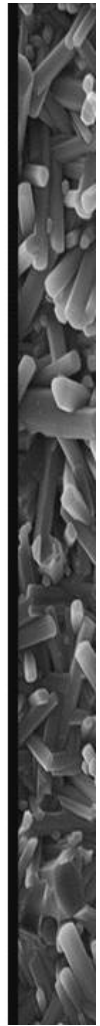


Composite: Cortical-Cancellous

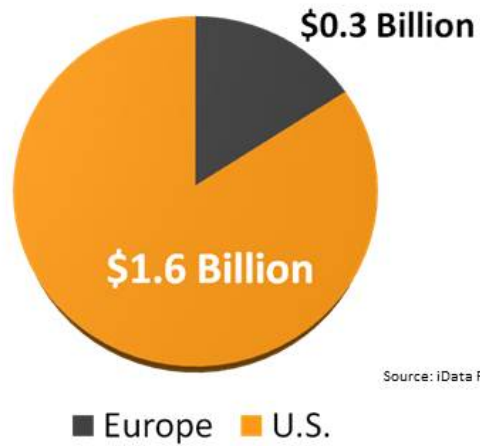
Synthetic bone for a variety of medical applications

Composite: Articular-Bone Ingrowth

Joint arthroplasty/resurfacing applications

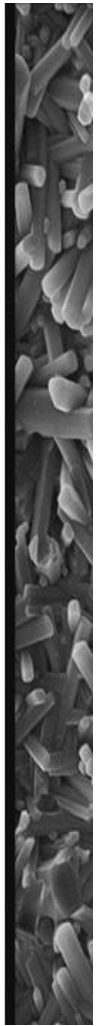


Sizable Spine Interbody Device Market



Source: iData Research Inc., 2014

- Sizable and growing market as surgeons look to replace PEEK
- Weaknesses of existing materials provide market share opportunities



Valeo™ Interbody Spinal Fusion Devices



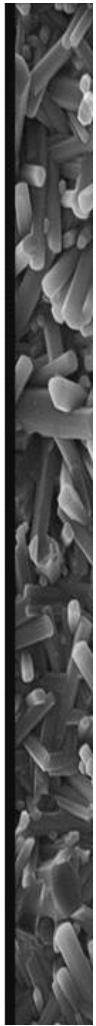
Full Line of FDA-cleared Si_3N_4 Products

- Cervical
- Lumbar
 - Anterior
 - Posterior
 - Oblique
 - Transforaminal
 - Lateral
- Corpectomy



Enhanced 2nd Generation Features

- Universal threaded instrumentation
- More implant sizes and approaches
- Improved safety & ease of use



Innovative Product Pipeline Key to Future Growth

Pipeline Products

- Composite cervical spinal fusion device
- Lateral lumbar product line expansion
- 2nd generation Si_3N_4
- Modular & cannulated pedicle screw system
- 3D printed Si_3N_4
- All-porous Si_3N_4 interbody
- Stand alone cervical interbody
- Si_3N_4 coatings on metals
- Brazing of Si_3N_4



Product Pipeline: Composite Spinal Fusion Device

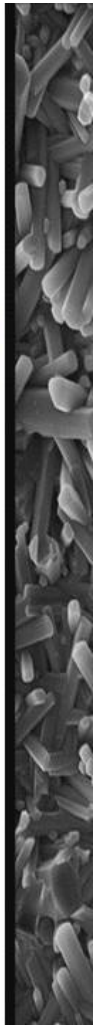


Pending FDA 510(k) Clearance

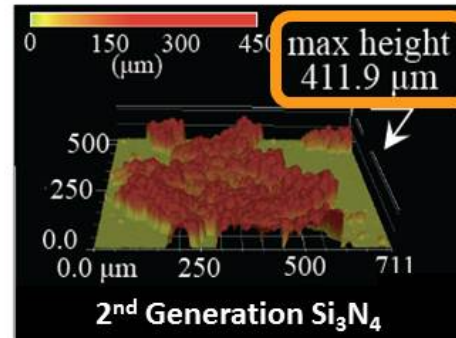
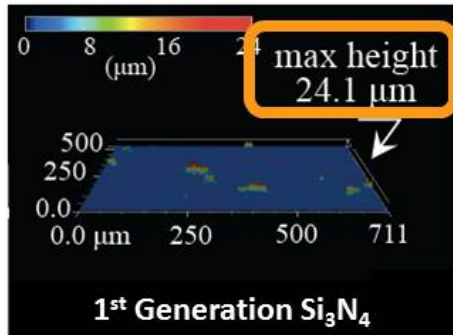
- Disruptive technology offering silicon nitride benefits
 - Super hydrophilic; orthobiologic alternative
 - Avoids possible use of cadaveric or autograft bone
 - Additional applications outside of spine

Silicon Nitride Statistically Equivalent to Industry Gold Standard

- Blinded, randomized clinical trial comparing Si_3N_4 to PEEK filled with autograft
- Statistically equivalent clinical improvement and fusion rates
- 24-month data submitted to FDA for anticipated composite device clearance



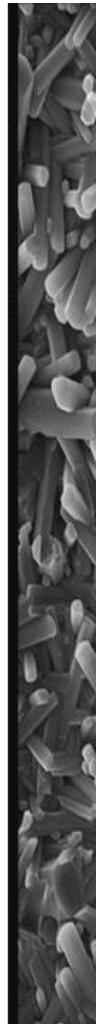
Product Pipeline: Next Generation of Silicon Nitride



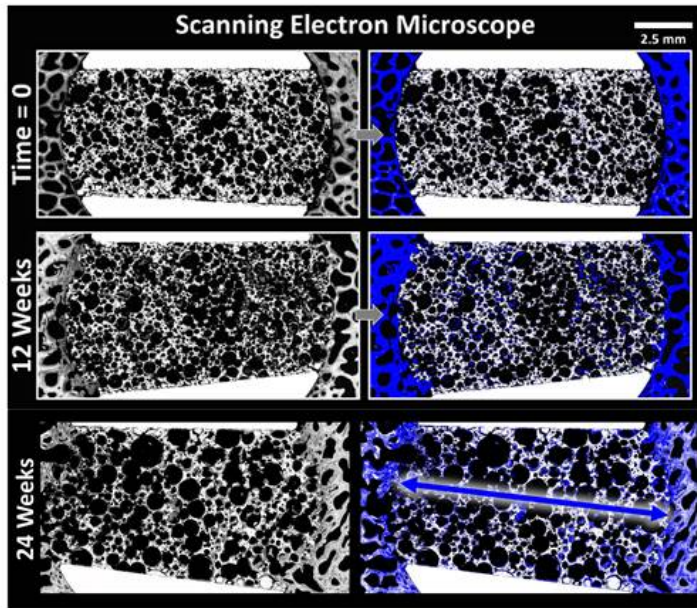
Exposing osteoblast-like Saos-2 cells* to biomaterial surfaces in an *in vitro* experiment demonstrates modulation of hydroxyapatite production rate

Osteoconductivity increased **190%** over current silicon nitride composition

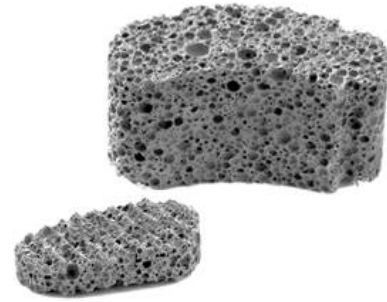
Pezzotti et al, "Silicon Nitride: A Synthetic Mineral for Vertebrate Biology," submitted to *Scientific Reports*
*Apatite deposition: 5×10^5 cell/mL, DMEM medium w/ 50 $\mu\text{g}/\text{mL}$ ascorbic acid, 10 mM β -glycerol phosphate, 100 nM dexamethasone, 10% FBS, 7 day incubation



Product Pipeline: Porous Fusion Device



In-vivo Ovine Medial Femoral Condyle Study



Bone In-growth = >3mm

Bone In-growth = 5.5mm

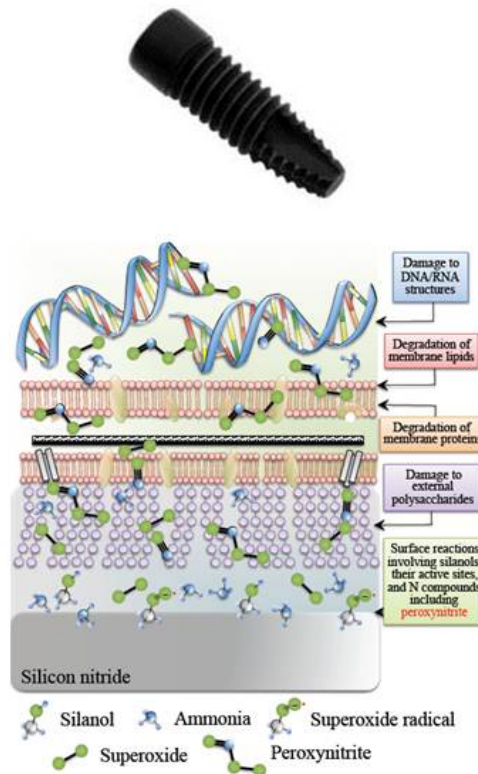
Product Pipeline: New Pedicle Screw System

- Modular
 - Can be implanted without the head – ability to distract of screws
 - Lends well to MIS system
- Tension head
 - Screw head is adjustable and will hold a specific angle or position
- Triple lead
 - Cortical trajectory approach
 - Aggressive option for those not using power
- Cannulated
 - Navigation and MIS friendly



Product Pipeline: Dental Implants

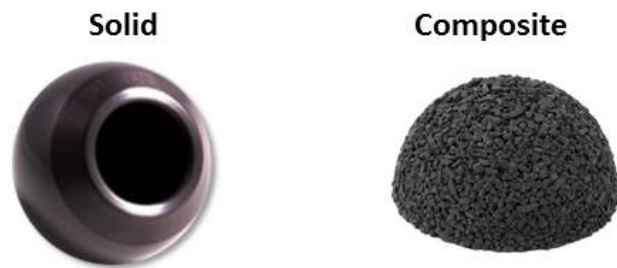
- Signed joint development agreement
 - Biomechanical and material testing to be completed in Q3 2016
- JDA to provide additional basis for FDA clearance process
- Recent peer-reviewed publication shows silicon nitride holds promise as a therapeutic aid for treating severe gum disease



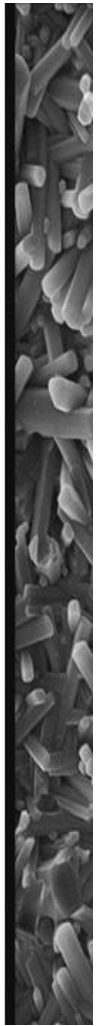
Langmuir, 2016, 32(12), pp 3024–3035

Product Pipeline: Hip & Knee Replacement

- No corrosion; no metal ion release
- Articulating and bone growth properties on a single device
- Additional testing¹ of the femoral head to further validate:
 - Reduced oxidization of poly surfaces
 - Superior burst strength and fracture toughness
 - Resistance to corrosion and fretting
 - Superior wear



¹ Femoral head testing to be done in partnership with Kyocera, Kyoto University and the University of Nebraska



Intellectual Property



60 patents issued

- 49 U.S.
- 11 International

30 patent applications

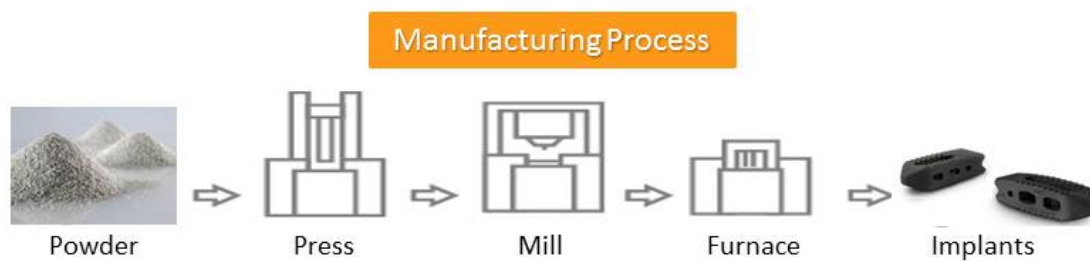
- 19 U.S.
- 11 International

Unique Manufacturing Technique

- 30,000 sq. ft. manufacturing facility in Salt Lake City, UT
 - Only FDA & CE cleared Si_3N_4 medical device manufacturing facility
 - Vertically integrated for rapid prototyping and development
 - Dedicated R&D and Product Development laboratories
- Production of powder and green compact preparation
- **Ability to manufacture complex designs and shapes**
- Rigorous quality control process for each implant



Silicon Nitride



Hybrid Sales Strategy

AMDA Spine

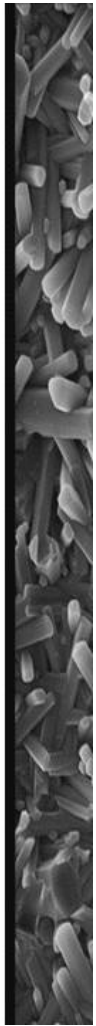
- Generate sales through independent domestic distributors
- Future growth through additional distributors & surgeons, while introducing new & innovative products
- Updating pedicle screw system to on-board additional new surgeons

Private Label

- Sell Amedica products with partner's logo and through their channels
- Near-term sales w/ limited selling expense and no CapEx
- Validates Si_3N_4 benefits **AND** penetrates market faster

OEM

- Convert existing partner's metal or plastic device to Si_3N_4
- Prototype, testing and FDA clearance in-house
- Longer-term revenue impact w/ limited additional spend
- Enhance body of scientific data to position Si_3N_4 as the standard of care



Key Partnerships

Current Partnerships

- Spinal Kinetics

- First private label and OEM partner



- Weigao Orthopedics

- 10 year exclusive distribution agreement with largest orthopedic company in China
- 6 years of minimum purchase requirements totaling 225,000 units (~10x total unit sales to-date)
- Finalizing CFDA clearance submission strategy



- Kyocera

- Secondary supplier source – Vancouver, WA



Potential Partnerships

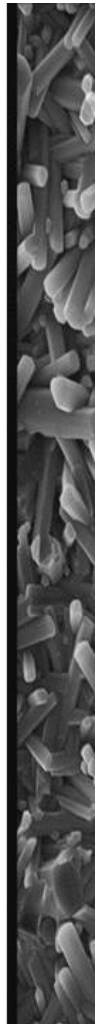
- Other Global Spine Partners

- Assessing regulatory clearance path with several potential partners in Japan, Taiwan and Australia

- Key Arthroplasty Discussions

- Current material testing agreements lead to clearer path toward additional OEM partners

- Non-medical Applications



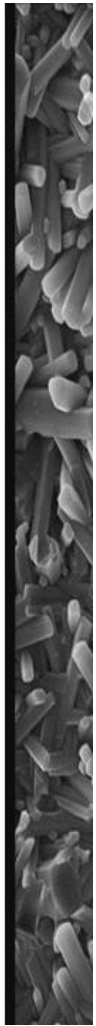
Business Highlights

2015

- Reduced principal debt balances from \$24.5 million to approx. \$18.0 million
- Received clearance for 1st generation S₃N₄ device and instrumentation in Brazil
- Signed four Private Label/OEM partnerships
- Entered the dental market with joint development agreement

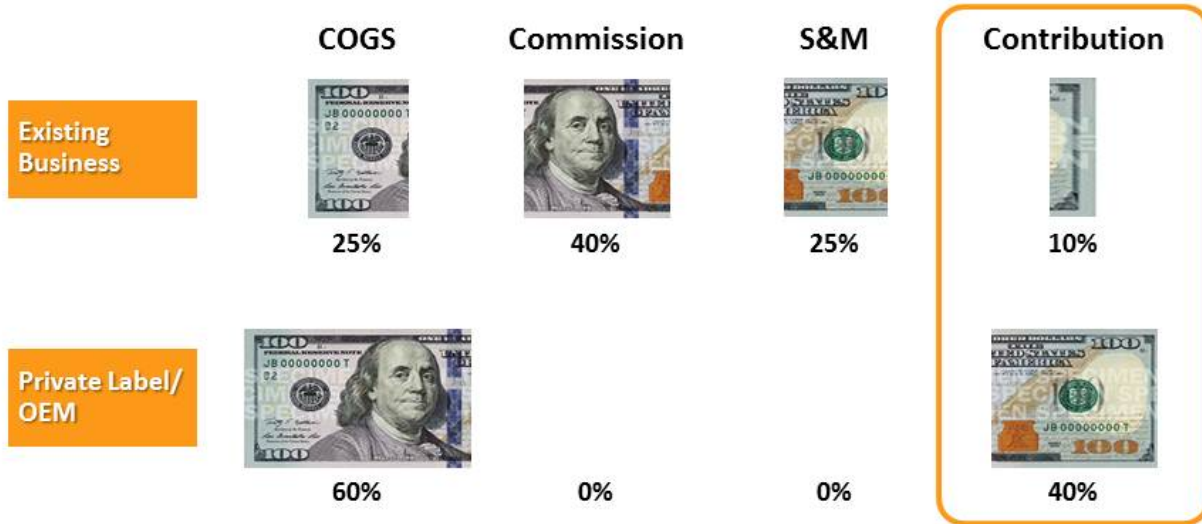
2016 Year-to-date

- Improved Q1 operational cash burn by 51%
- Signed an exclusive silicon nitride distribution agreement with the largest Chinese orthopedic company
- Completed debt exchange agreement and further reduced debt by 32% to \$12.1 million
- Met in-person with the FDA regarding the Company's composite silicon nitride fusion device
- Became first to 3D print silicon nitride ceramic for medical applications
- Featured unique silicon nitride attributes in several key industry events and publications
- Launched Valeo II™ silicon nitride cervical interbody fusion system
- Signed additional material testing agreements with key orthopedic, dental and other companies
- Announced collaboration on biologically enhanced spinal fusion devices



Operating Contribution Margin by Department

Before overhead allocation



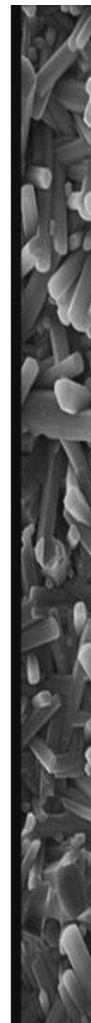
Financial Summary

- **Q1 2016 results:**

- Total revenue of \$4.2 million
- Gross margins increased to 79% from 68% year-over-year
- Operating expenses decreased by 20% year-over-year
- Operational cash burn decreased by \$1.5 million or 51% year-over-year

- **2015 results**

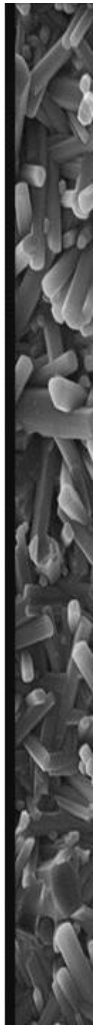
- Total revenue of \$19.5 million
- Cash used in operations down \$5.5 million or 38% year-over-year
- Principal debt balances reduced from \$24.5 million to \$18.0 million



Capitalization Table

Common Shares Outstanding (as of 6-6-16)	13,306,000
Warrants Outstanding	807,000
Unit Option	38,000
Options Outstanding (as of 3-31-16)	122,000
# Shares for Convertible Note (if converted at 6-6-16)	489,000
Total Potentially Dilutive Securities	1,456,000
Total Shares & Potentially Dilutive Securities	14,762,000

Hercules Term Loan	\$10,600,000
Magna Term Loan	\$800,000
Convertible Note	\$700,000
Total Debt Outstanding (as of 6-6-16)	\$12,100,000



Thank You

Nasdaq: AMDA

Investor Relations

Mike Houston
IR@amedica.com
801-839-3534

